

Effects of serum proteins on corrosion behavior of ISO 5832–9 alloy modified by titania coatings

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Abstract Stainless steel ISO 5832–9 type is often used to perform implants which operate in protein-containing physiological environments. The interaction between proteins and surface of the implant may affect its corrosive properties. The aim of this work was to study the effect of selected serum proteins (albumin and γ -globulins) on the corrosion of ISO 5832–9 alloy (trade name M30NW) which surface was modified by titania coatings. These coatings were obtained by sol–gel method and heated at temperatures of 400 and 800 °C. To evaluate the effect of the proteins, the corrosion tests were performed with and without the addition of proteins with concentration of 1 g L⁻¹ to the physiological saline solution (0.9 % NaCl, pH 7.4) at 37 °C. The tests were carried out within 7 days. The following electrochemical methods were used: open circuit potential, linear polarization resistance, and electrochemical impedance spectroscopy. In addition, surface analysis by optical microscopy and X-ray photoelectron spectroscopy (XPS) method was done at the end of weekly corrosion tests. The results of corrosion tests showed that M30NW alloy both uncoated and modified with titania coatings exhibits a very good corrosion resistance during weekly exposition to corrosion medium. The best corrosion resistance in 0.9 % NaCl solution is shown by alloy samples modified by titania coating annealed at 400 °C. The serum proteins have no significant effect onto corrosion of investigated biomedical steel. The XPS results confirmed the presence of proteins on the alloy surface after 7 days of immersion in protein-containing solutions.

Keywords Corrosion · Corrosion rate · Biomedical steel · Serum proteins · Sol–gel coating · Titanium dioxide

Introduction

The term “biomaterial” was defined by Williams (1987) as any substance (other than a drug) or combination of substances, synthetic or natural in origin, which can be used for any period of time as a whole or as a part of a system which treats, augments, or replaces any tissue, organ, or function of the body [1]. Metals, polymers, ceramics, and composites may be used as biomaterials. Each type of these materials has specific properties which are needed in a specific application field. Metallic biomaterials find an application as prostheses and implants in many medical disciplines, e.g., in orthopedics, cardiology, and dentistry. All of the requirements set out for metallic biomaterials are detailed in International Standard ISO 5832. This standard contains several parts concerned with titanium-based materials, cobalt-based materials, and stainless steels. Part 9 of the standard describes the chemical composition and physicochemical properties of wrought high nitrogen stainless steel dedicated for orthopedic implants [2]. The main alloying elements of this group of steel are as follows (wt%): iron (base), chromium (19.5–22.0), nickel (9.0–11.0), manganese (2.0–4.45), molybdenum (2.0–3.0), and nitrogen (0.25–0.5).

An ideal metallic biomaterial is expected to exhibit a very high biocompatibility, good corrosion resistance, and good mechanical properties. It is very difficult to combine all these properties in only one material. Improvement of mechanical properties, corrosion resistance, and biocompatibility of biomedical steels for implantology may be achieved through surface modification, for example by surface coating. A very popular material for biomedical coating is titanium dioxide (titania, TiO₂) due to its excellent corrosion, mechanical, and biological properties [3, 4]. Many methods are used to prepare biomedical coatings. One of them is sol–gel method [4–7].

Orthopedic implants, made of stainless steel ISO 5832–9 type, operate in protein-containing physiological environments. Blood plasma contains three types of dissolved proteins—

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albumin, globulins, and fibrinogen. Following the implantation of a biomaterial, the first event to occur at the tissue-material interface is protein adsorption [8, 9]. The layer of adsorbed proteins is formed immediately after metal immersion in human plasma [10]. The interaction between proteins and surface of the implant may affect its corrosive properties. Conclusions concerning the influence of proteins are not clear. Some authors [11, 12] report that bovine serum albumin (BSA) diminishes the corrosion rate of stainless steels and titanium alloys, while other authors [13, 14] report that BSA promotes the dissolution of these alloys by forming stable complexes with the metallic components of the alloys. Literature data show that the effect depends on the type of biomaterial [15] and its surface finish [16], as well as on the concentration of protein [15] and pH of the solution [17].

The aim of this work was to study the effect of selected serum proteins (albumin and γ -globulins) added to the physiological saline solution (0.9 % NaCl, pH 7.4) on the corrosion of ISO 5832–9 alloy (trade name M30NW) which surface was modified by titania coatings.

Experimental

Materials

The biomedical alloy M30NW (Aubert & Duval, France) was used as a metallic substrate. Its chemical composition is as follows (wt%): Cr (20.62), Ni (9.53), Mn (4.13), Mo (2.12), N (0.40), Nb (0.31), C (0.032), and Fe (rest). M30NW alloy samples were discs with a diameter of 22 mm and a thickness of about 3 mm. Every sample was ground on SiC abrasive paper (80–1,500 grits), polished with Al₂O₃ suspension (0.3 μ m), chemically etched in a mixture of 2 % HF, 10 % HNO₃, and 88 % H₂O, passivated in boiling water (10 min), rinsed with ethanol, and dried with argon. Samples prepared in such a way were ready for TiO₂ coating. The precursor for TiO₂ coating was titanium (IV) isopropoxide (Ti[OCH(CH₃)₂]₄, 97 %, Aldrich) [7]. Titania coating (as a single layer) was applied onto alloy surface by dip-coating technique using a dip-coater DCMono 75 (NIMA Technology). The final annealing of applied coating was carried out in a muffle furnace at temperatures of 400 and 800 °C for 1 h.

All the chemical reagents used in the experiment were of analytical grade and were applied without further purification.

Corrosion medium

The corrosion medium was composed of sodium chloride (NaCl, pure p.a., POCH S.A.), bovine serum albumin (BSA, >96 %, Sigma), γ -globulins from bovine blood (\geq 97 %, Sigma), and distilled water. The starting composition in 1,000 mL of each solution and their codes are listed in

Table 1. All solutions were buffered by phosphate buffer (KH₂PO₄/Na₂HPO₄ × 12H₂O) and adjusted to pH 7.4 using hydrochloric acid (HCl, pure p.a., POCH S.A.).

Electrochemical measurement

During the weekly corrosion tests, all samples immersed in corrosion medium were stored at constant temperature of 37 °C. On selected days (0, 1, 3, 7), non-destructive corrosion tests were performed using potentiostat/galvanostat PGSTAT 128 with FRA2 module (EcoChemie Autolab). The corrosion measurements were carried out in three-electrode electrolytic cell with a sample as a working electrode, platinum foil with an area of 60 cm² as a counter electrode, and saturated calomel electrode (SCE, $E^\circ=0.241$ V vs SHE) as a reference electrode. All potentials reported in this paper are given versus SHE electrode. Exposed area of each sample was approximately 0.64 cm².

Some basic corrosion parameters such as corrosion potential, polarization resistance, and electrochemical impedance, were determined versus immersion time. The corrosion potential (E_{cor}) as open circuit potential was measured for 2,000 s. The polarization resistance (R_p) of the specimen in the test solution was measured using linear polarization method. The measurements were done in a scanning range of ± 20 mV versus E_{cor} potential with a scan rate of 0.3 mV s⁻¹. Electrochemical impedance characteristics were registered at E_{cor} potential by applying a sinusoidal signal of ± 10 mV within the frequency range from 0.01 to 10 kHz. Each measurement was maintained at 37 °C. Three specimens were tested for each corrosion test and results presented in this paper are averaged values.

Surface characterization

The specimens were removed from the solutions after immersion for 7 days and rinsed with distilled water. Next, their surfaces were examined by optical microscopy and X-ray photoelectron spectroscopy (XPS). The optical images were obtained using metallographic microscope MMT 800BT (Mikrolab, Poland). All XPS spectra were recorded with a PHI 5000 VersaProbe scanning spectrometer (ULVAC-PHI, Japan/USA). The X-ray source used was monochromatic Al-K α radiation (1486.6 eV) with an input power of 25 W. X-ray beam was focused to a diameter of 100 μ m; measured area was defined as 1,000 μ m × 1,000 μ m. The electron energy analyzer was operated at a pass energy of 117.4 eV (energy step 0.4 eV) for fast survey scan, and 23.5 eV (energy step 0.1 eV) for all high-resolution spectra. All measurements were conducted with the use of a charge neutralizer system. The angle between sample surface and electron energy analyzer was 45°. The binding energy scale was normalized to the C1s peak (285.0 eV).

Table 1 Values of corrosion rate (mm year⁻¹) at 0 and 7th day of immersion

Sample	Uncoated M30NW		M30NW+TiO ₂ 400		M30NW+TiO ₂ 800	
	0	7	0	7	0	7
NaCl	3.97×10^{-5}	1.37×10^{-5}	6.40×10^{-6}	5.11×10^{-6}	2.80×10^{-5}	1.29×10^{-5}
BSA	6.00×10^{-5}	1.27×10^{-5}	4.94×10^{-6}	5.03×10^{-6}	3.72×10^{-5}	1.13×10^{-5}
gG	4.98×10^{-5}	9.16×10^{-6}	5.57×10^{-6}	5.83×10^{-6}	3.74×10^{-5}	1.78×10^{-5}

Standard deviation of CR was less than 10 %

Results

Electrochemical measurements

Figure 1 presents the results of corrosion potential E_{cor} versus immersion time. For uncoated samples (Fig. 1a) in solution without proteins, the E_{cor} value is rising close to 0.23 V_{SHE} at

the 7th day. The positive potential evolution indicates the formation of a protecting passivation layer on the M30NW alloy. Such evolution of E_{cor} values is not observed in the protein-containing solutions. In case of samples modified by titania coatings (Fig. 1b, c) a regression with time of the E_{cor} is observed. This regression is similar in solutions both with and without proteins.

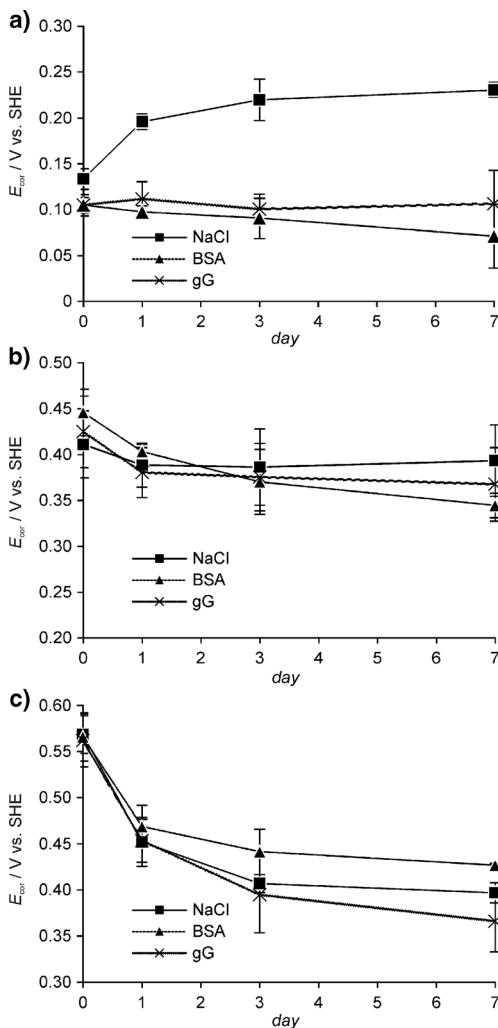


Fig. 1 Corrosion potential E_{cor} versus immersion time for uncoated alloy (a), alloy with TiO₂ coating annealed at 400 °C (b), and alloy with TiO₂ coating annealed at 800 °C (c)

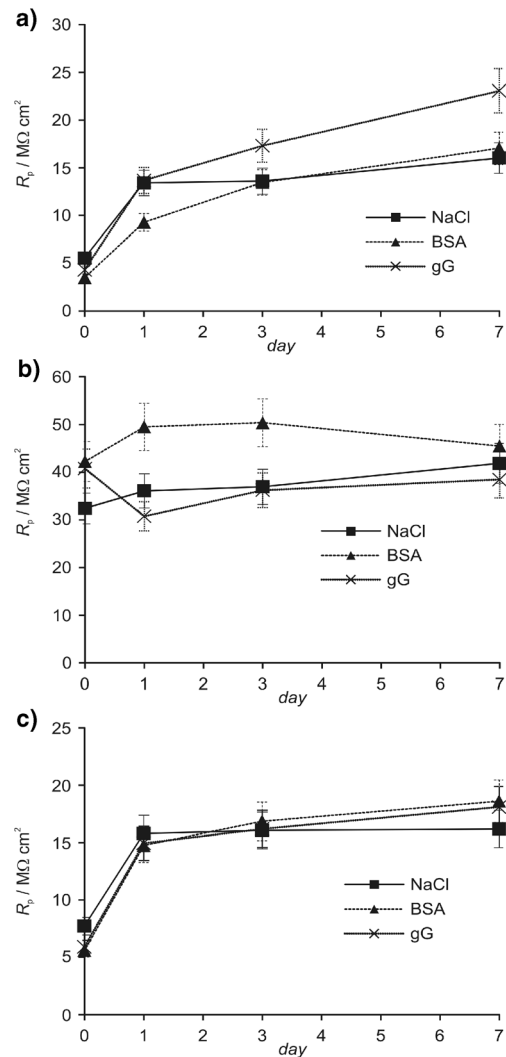


Fig. 2 Polarization resistance R_p versus immersion time for uncoated alloy (a), alloy with TiO₂ coating annealed at 400 °C (b), and alloy with TiO₂ coating annealed at 800 °C (c)

The values of polarization resistance were determined from the slope of the linear polarization characteristics using CorrView software (Scribner Associates Inc.). Figure 2 presents the polarization resistance-immersion time dependence for all investigated samples. It can be seen that the longer the immersion time, the higher the value of R_p in case of both uncoated samples (Fig. 2a) and samples coated with TiO_2 annealed at 800°C

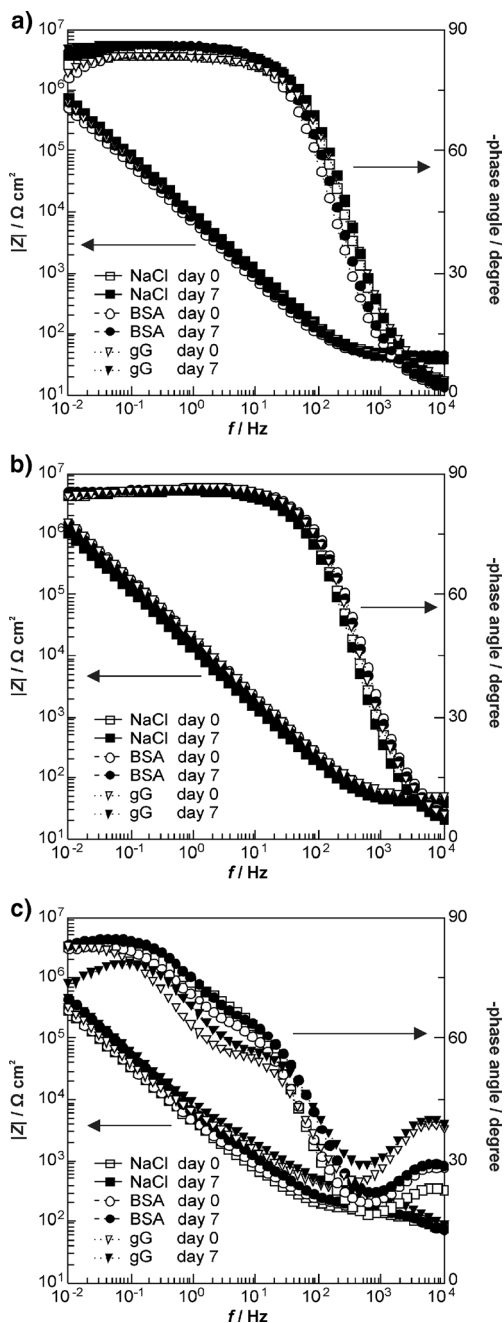


Fig. 3 Impedance spectra (Bode plots) recorded on 0 and 7th day of immersion of uncoated alloy (a), alloy with TiO_2 coating annealed at 400°C (b), and alloy with TiO_2 coating annealed at 800°C (c) in all corrosion solutions

(Fig. 2c). In case of samples with titania coating annealed at 400°C (Fig. 2b), the R_p values are relatively stable during the immersion time.

Collected values of polarization resistance were used for determination one of the most important corrosion parameters—corrosion rate. The values of corrosion rate CR were calculated from the Eq. (1) according to procedure described by ASTM Standard G 102–89 [18], which assumes that at E_{cor} potential, only uniform type of corrosion is occurring:

$$CR = K_1 \frac{i_{\text{cor}}}{\rho} EW \quad (1)$$

where CR is the corrosion rate (mm year^{-1}), $K_1 = 3.27 \cdot 10^3 \text{ mm g} (\mu\text{A cm year})^{-1}$, i_{cor} is the corrosion current density ($\mu\text{A cm}^{-2}$) calculated on the basis of linear polarization characteristics, ρ is the density of investigated alloy which is 7.90 g cm^{-3} , and EW is the equivalent weight which for M30NW alloy equals to 19.18 in the experimental set used in this study.

To calculate the CR value, it was assumed that at corrosion potential, only the substrate, M30NW alloy, corrodes. No other corrosion processes associated with the formed and annealed TiO_2 coatings were taken into account in this calculation. According to the ASTM standard [18], only elements above one mass percent in the alloy were included in EW calculation.

Table 1 presents the values of corrosion rate at 0 and 7th day of immersion. In case of uncoated samples and samples coated with TiO_2 annealed at 800°C , a decrease in corrosion rate can be observed in all solutions while immersion time increases, but this behavior is more pronounced in the protein-containing solutions. The values of corrosion rate of samples coated with TiO_2 annealed at 400°C are very low and they are independent on the presence of serum proteins.

Impedance spectra for all investigated samples recorded at the beginning and the end of immersion tests are presented in Fig. 3. Impedance spectra for uncoated alloy (Fig. 3a) show an increase of impedance during 7 days of immersion. This increase seems to be greater in case of protein-containing

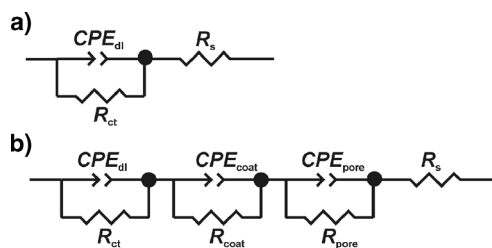


Fig. 4 The equivalent circuits used for quantitative evaluation of impedance spectra for uncoated alloy and alloy with TiO_2 coating annealed at 400°C (a), and samples with TiO_2 coating annealed at 800°C (b)

Table 2 Values of the parameters obtained from the fitting procedure at 0 and 7th day of immersion

Sample	Day	$R_s/\Omega\text{ cm}^2$	CPE_{dl}		$R_{ct}/M\Omega\text{ cm}^2$	CPE_{pore}		$R_{pore}/\Omega\text{ cm}^2$	CPE_{coat}		$R_{coat}/\Omega\text{ cm}^2$	
			$C_{dl}/\mu\text{F cm}^{-2}$	n		$C_{pore}/\mu\text{F cm}^{-2}$	n		$C_{coat}/\mu\text{F cm}^{-2}$	n		
Uncoated alloy												
NaCl	0	42.7	21.4	0.93	19.5	–	–	–	–	–	–	
	7	40.9	18.6	0.94	58.6	–	–	–	–	–	–	
BSA	0	46.3	25.9	0.95	4.2	–	–	–	–	–	–	
	7	45.6	21.5	0.95	54.2	–	–	–	–	–	–	
gG	0	37.7	22.1	0.92	11.9	–	–	–	–	–	–	
	7	37.4	19.5	0.95	115.4	–	–	–	–	–	–	
With TiO ₂ 400 °C												
NaCl	0	44.3	11.4	0.95	59.1	–	–	–	–	–	–	
	7	41.5	13.5	0.95	80.8	–	–	–	–	–	–	
BSA	0	43.8	8.9	0.96	58.2	–	–	–	–	–	–	
	7	41.5	10.2	0.95	221.1	–	–	–	–	–	–	
gG	0	49.4	9.0	0.96	40.5	–	–	–	–	–	–	
	7	46.2	13.1	0.95	114.1	–	–	–	–	–	–	
With TiO ₂ 800 °C												
NaCl	0	47.7	44.8	0.94	11.4	4.9	0.76	104.1	172.9	0.71	1446	
	7	46.4	28.4	0.95	12.6	5.0	0.76	111.2	129.1	0.73	1206	
BSA	0	39.3	45.4	0.94	10.0	4.4	0.76	128.5	108.7	0.74	1544	
	7	38.0	29.4	0.95	13.2	4.4	0.77	125.2	79.6	0.75	1442	
gG	0	39.0	40.7	0.93	51.7	2.2	0.81	220.9	53.3	0.72	3270	
	7	36.0	25.3	0.90	3.3	2.5	0.80	222.1	38.8	0.75	2183	

solutions. In case of alloy samples with TiO₂ coating annealed at 400 °C (Fig. 3b), their impedance do not depend on immersion time and solution chemistry—the impedance spectra are almost identical at day 0 and at day 7 for all solutions. The samples with titania coating heat-treated at 800 °C (Fig. 3c) have different impedance characteristics—the Bode plots show three time constants in all corrosion solutions.

All obtained impedance data were fitted by the equivalent electrical circuits using Zview2 software. The quality of the fitting was judged by the χ^2 value that was lower than 4×10^{-4} . Instead of pure capacitors, constant phase elements (CPE) were introduced in the fitting procedure to obtain good agreement between the

experimental and simulated data. The impedance of CPE is defined by the Eq. (2):

$$Z_{CPE} = \frac{1}{Q(j\omega)^n} \tag{2}$$

where Q ($\Omega^{-1} \text{ s}^n \text{ cm}^{-2}$) is the combination of properties related to both the surface and the electroactive species independent on frequency, n is related to a slope of the $\log Z$ versus $\log f$ (Bode plot), and ω is the angular frequency.

In the ideal case when the exponential factor $n=1$, the CPE acts as a pure capacitor. In practice, n deviates from 1 so the fitting results are given as both the parameter Q and the exponent “ n .” It should be noted that for simplicity, Q is often considered as a capacitance [19].

For uncoated alloy and alloy with TiO₂ coating annealed at 400 °C, the best data-fitting equivalent circuit (Fig. 4a) is characterized by parallel combination term (R, CPE) in series with the resistor R_s . The values of the parameters obtained with the fitting procedure versus immersion time are reported in Table 2. Resistor R_s determines uncompensated electrolyte resistance and it is about $40\Omega\text{ cm}^2$ for all types of corrosion medium. Constant phase element CPE_{dl} is equivalent to double layer capacity (since the factor n has a value in the range of 0.92–0.96, CPE can be said to behave similarly to a pure

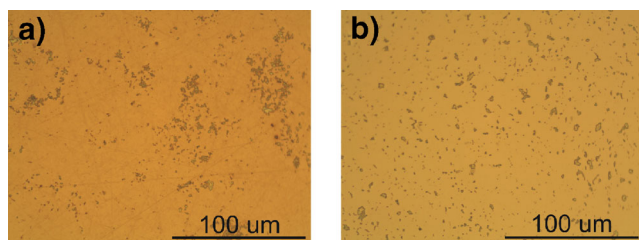


Fig. 5 Optical microscope images for M30NW alloy with TiO₂ coating annealed at 400 °C obtained after 7 days of immersion in protein-containing solutions: BSA (a) and gG (b)

capacitor). Resistor R_{ct} reaches a value in the order of $10^7 \Omega \text{ cm}^2$ and it is related to charge transfer resistance of corrosion processes. It is in agreement with R_p value obtained from linear polarization characteristics.

In case of alloy samples coated by TiO_2 layer annealed at 800°C , the simple equivalent circuit (Fig. 4a) was inadequate; therefore, a more complex circuit (Fig. 4b) has been used. This

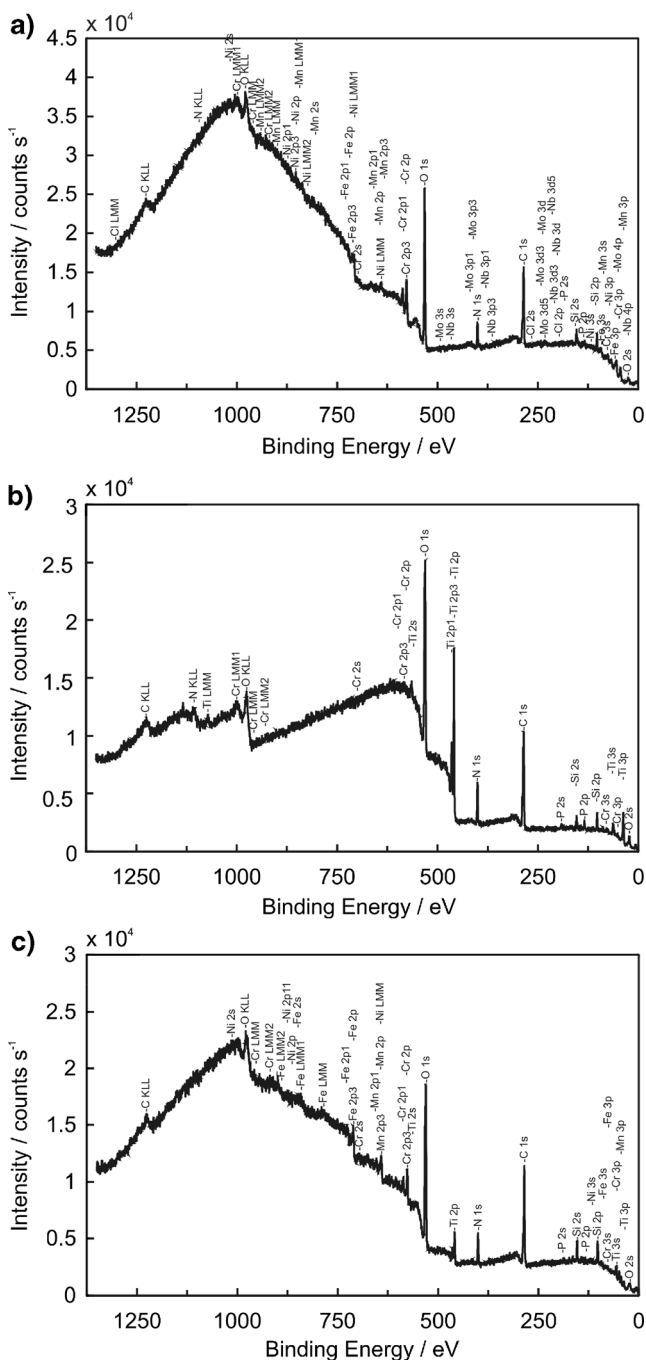


Fig. 6 The representative survey spectra for samples immersed in BSA-containing solution: uncoated sample (a), sample with TiO_2 annealed at 400°C (b), and samples with TiO_2 annealed at 800°C (c)

circuit contains three parallel combination terms (R , CPE) since three time constants are present on the Bode plots. The differences result from entirely different surface properties of TiO_2 coating depending on applied annealing temperature. In previous paper [7], we have reported that TiO_2 sol–gel coating heat-treated at 800°C is a porous, diffusive layer which contains new TiO_2 -based phases instead of pure TiO_2 crystal structures. Therefore, in addition to solution resistance R_s , double layer capacitance CPE_{dl} , and charge transfer resistance R_{ct} , the electrical equivalent circuit contains elements associated with pores and coating properties. The values of the parameters obtained with the fitting procedure versus immersion time are reported in Table 2. Constant phase element CPE_{pore} and resistor R_{pore} describe porosity of coating layer. These elements are related with time constant visible in high frequency range in Bode plots (Fig. 3c). Low R_{pore} values (in the order of hundreds of $\Omega \text{ cm}^2$) are to be expected since they correspond to the low electrolyte resistance inside the pores [19]. Constant phase element CPE_{coat} and resistor R_{coat} characterize surface properties of TiO_2 layer. These elements are related with time constant visible in medium frequency range in Bode plots (Fig. 3c). The low n values obtained for CPE elements are typical for rough surface of the coating [20]. The rough surface is a result of dislocations, grain boundaries, and precipitations. CPE element is a practical way to describe the above interfacial characteristics [21].

Surface analysis after 7 days of immersion

At the end of the corrosion tests, the alloy samples were observed by optical microscope in order to analyze the changes in surface morphology. Optical microscope results did not reveal any local corrosion damage on the investigated samples. This fact indicate that during weekly corrosion tests, only general corrosion occurred. Additionally, for all samples immersed in solutions with proteins, some sediment on the surface was observed. The representative microscope images obtained for alloy samples with TiO_2 coating annealed at 400°C are presented in Fig. 5.

This sediment may be related to presence of proteins adsorbed on the sample surfaces. In order to clarify its origin, the surface analysis was performed using XPS method. X-ray spectra in a wide range of binding energy were collected for both uncoated samples and samples with titania coating after 7 days of immersion in corrosion medium. Figure 6 presents a representative XPS survey spectra taking each type of samples after immersion in BSA solution for example. Surface film composition of all alloy samples after 7 days exposure to the corrosion medium is presented in Table 3. The general features of the spectra are peaks of carbon (C1s) and oxygen (O1s) which are observed on all the samples. Carbon concentration is ranged from 16 to 57 at.%. The XPS results exhibit the presence of main alloying components (Fe, Cr, Ni, Mn, Mo) in case

Table 3 Surface film composition of M30NW alloy after 7 days exposure to the corrosion medium

Sample/Solution	Atomic concentration (at.%)										
	Ni 2p	Fe 2p	Mn 2p	Cr 2p	O 1s	Ti 2p	N 1s	C 1s	Mo 3d	Nb 3d	P+Si 2p
Uncoated alloy											
NaCl	0.50	0.27	0.12	6.05	34.01	–	1.65	50.76	0.23	0.00	6.40
BSA	0.50	0.63	0.00	5.78	37.51	–	6.25	41.17	0.22	0.06	7.89
gG	0.05	0.08	0.00	0.90	23.07	–	12.77	57.70	0.08	0.00	5.34
With TiO ₂ 400 °C											
NaCl	–	0.64	–	2.19	53.97	18.54	2.38	20.16	–	–	2.14
BSA	–	0.51	–	1.21	43.02	12.07	7.15	36.03	–	–	0.00
gG	–	0.15	–	0.45	26.64	4.09	12.47	56.18	–	–	0.00
With TiO ₂ 800 °C											
NaCl	–	2.66	4.19	5.03	46.34	3.64	2.73	28.41	–	–	7.00
BSA	–	1.87	2.28	2.65	37.12	2.58	5.55	40.12	–	–	7.87
gG	–	0.52	0.80	1.26	28.59	1.20	9.62	50.94	–	–	7.07

of uncoated samples. Some of the alloying elements (Fe, Cr, Mn) are also visible on the surface of alloy samples coated with TiO₂ layer annealed at 800 °C. In case of samples modified with TiO₂ coating annealed at 400 °C, only signals of Fe and Cr coming from the substrate were stated. Additionally, signals of Ti and O coming from the titania coating were stated for both modified alloy samples. The XPS results show also that nitrogen (N1s) is present on all the samples. In case of samples exposed to solution without addition of protein, the atomic concentration of N is lower than 3 %. Much higher percentage of nitrogen is observed for samples immersed in protein-containing solutions. Approximately two times as much nitrogen was found on the surface of samples after contact with a solution containing γ -globulins than for samples immersed in albumin-containing solution. It should be noticed that the higher the amount of nitrogen, the less the amount of iron and chromium found on the alloy's surface. An example of high-resolution spectra for N1s region which were used for detection of nitrogen percentages is presented in Fig. 7. Percentages of N

atoms on surfaces after immersion in protein-containing solutions obtained by XPS analysis are summarized in Fig. 8.

Discussion

The results of corrosion tests presented in this study confirm that titania sol–gel coatings improve the corrosion resistance of M30NW biomedical alloy in 0.9 % NaCl solution. This improvement is dependent on the annealing temperature used. In our previous paper [7], we characterized the structure and character of interactions between substrate and deposited TiO₂ sol–gel layers in relation to different annealing temperatures. We have stated that the coating annealed at temperature of 400 °C has an adhesive character and it has an amorphous structure with anatase crystallites. Whereas, TiO₂ coating annealed at 800 °C has a diffuse character and contains new phases (NiTiO₃ and Fe₂TiO₄) due to the interaction with steel

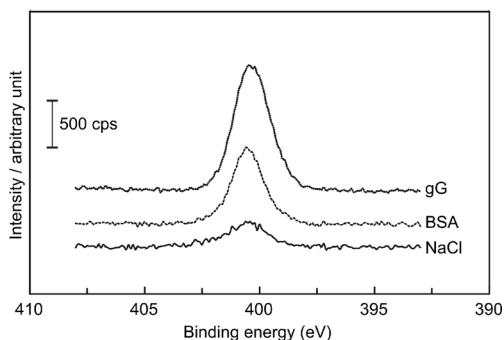


Fig. 7 The XPS high-resolution N1s spectra for M30NW alloy with TiO₂ coating annealed at 400 °C (the curves are shifted by 300 counts relative to each other for better separation)

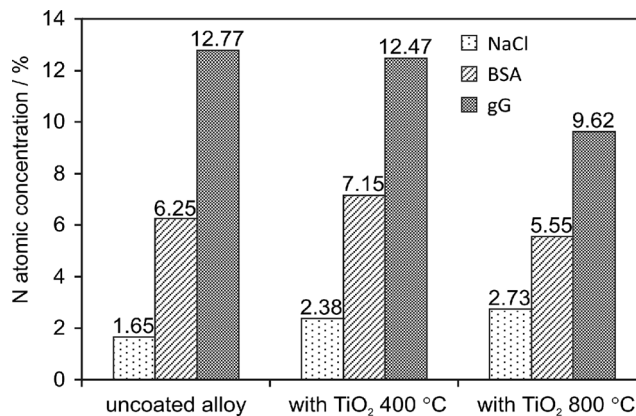


Fig. 8 Percentages of N atoms on surfaces after immersion in protein-containing solutions obtained by XPS

substrate. Such differences in crystalline structure and character of substrate layer interaction imply differences in anticorrosion properties of titania coatings [7].

The results presented in this paper indicate that M30NW alloy both uncoated and modified with titania coatings has a very good corrosion resistance also during weekly exposition to corrosion medium with and without addition of serum proteins. The results of corrosion potential versus immersion time show a very weak interaction of proteins with surfaces modified by titania coatings, while in case of unmodified alloy, it is significant. The polarization resistance of uncoated samples and samples with titania coating obtained at 800 °C increases, and corrosion rate decreases during exposure to all corrosion solutions. The samples with titania coating obtained at 400 °C show very good corrosion resistance in all solutions—polarization resistance is in the range of 30–50 M Ω cm² not depending on the presence of proteins and immersion time. Results of impedance measurements did not show any significant effect of proteins onto weekly corrosion of investigated biomedical steel. However, they indicate the differences in phase boundary electrolyte solution/TiO₂-coated alloy depending on annealing temperature.

The XPS results allowed to determine the changes in surface composition of analyzed samples after 7 days of exposure to the protein-containing solutions. Proteins (albumins and γ -globulins) as chemical molecules have carboxyl (–COOH), amino (–NH₂), and amido (–CONH–) groups. Hence, the major elements of the proteins are oxygen, carbon, and nitrogen. Carbon contamination is unavoidable in XPS studies; therefore, carbon signal cannot act as a protein indicator. We assumed that the nitrogen signal is indicative of the presence of a protein. This assumption is in accordance with Dupont-Gillain et al., Feng et al., and Karimi et al. [9, 22, 23]. The results of surface analysis obtained by X-ray photoelectron spectroscopy clearly showed that contact of samples with protein-containing solutions per 7 days causes increase in percentages of N atoms on their surfaces. This fact confirms the presence of proteins on the alloy surface after 7 days of immersion in protein-containing solutions.

Some researchers have suggested that the electrostatic attraction between proteins and biomaterials is important [13, 24, 25]. The isoelectric point of BSA is in the range of 4.2–5.5 [13, 22, 26], and in case of γ -globulins, it is in the range of 6.5–7.0 [26] so both proteins are negatively charged in the solution at pH 7.4. The pH zero point of charge for Fe-oxide/hydroxide, which is the outer oxide layer on a stainless steel surface, is 8.5 [13]. Hence, at pH 7.4, the stainless steel surface at the E_{cor} potential (Fig. 1a) has a net positive charge. This will facilitate the involvement of negatively charged carboxylate groups of the proteins as anchoring sites in the contact region between the proteins and the stainless steel surface [13]. However, for the interaction between proteins used in this study and TiO₂ coating, the electrostatic effects should not

be dominant factors since bovine serum albumin, γ -globulins, and TiO₂ (isoelectric point in the range of 4.7–6.2 [20]) are negatively charged in the solution at pH 7.4. Therefore, in our experimental set, a chemical adsorption of both proteins onto alloy surface modified with titania coatings is more probable rather than a physisorption. The chemical adsorption of BSA onto titanium surfaces through the interaction of C and O groups with the surface hydroxyl groups was reported by Feng et al. [22].

Conclusions

The corrosion of ISO 5832–9 alloy (trade name M30NW) which surface was modified by titania sol–gel coatings was investigated in this study using 0.9 % NaCl solutions with addition of two serum proteins (albumin and γ -globulins). The experimental results have led us to the following conclusions:

- M30NW alloy both uncoated and modified with titania coatings has a very good corrosion resistance during weekly exposition to corrosion medium. The best corrosion resistance in 0.9 % NaCl solution is shown by alloy samples modified by titania coating annealed at 400 °C.
- The serum proteins used in this study have no significant effect onto corrosion of investigated biomedical steel.
- The results of surface analysis by X-ray photoelectron spectroscopy clearly showed that contact of samples with protein-containing solutions per 7 days causes increase in percentages of N atoms on their surfaces. This fact confirms the presence of proteins on the alloy surface after 7 days of immersion in protein-containing solutions.
- In case of uncoated alloy, a physisorption probably occurs, while in case of alloy modified with titania coatings, a chemical adsorption is more probable.

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